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Publication number : **0 569 122 A1**

⑫

## EUROPEAN PATENT APPLICATION

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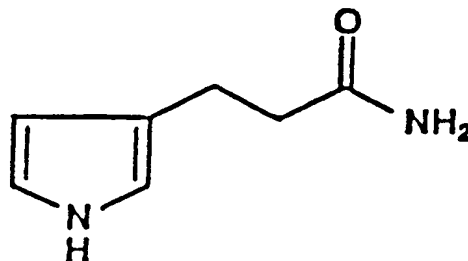
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⑫⑤④ Pyrrole propionic acid amide, its production and use as calpain inhibitor.

⑫⑤⑦ A compound of formula :



(1)

or a pharmaceutically or veterinarily acceptable salt thereof, is useful as a calpain inhibitor.

EP 0 569 122 A1

This invention relates to a novel calpain inhibitor, cystamidin A and to its production.

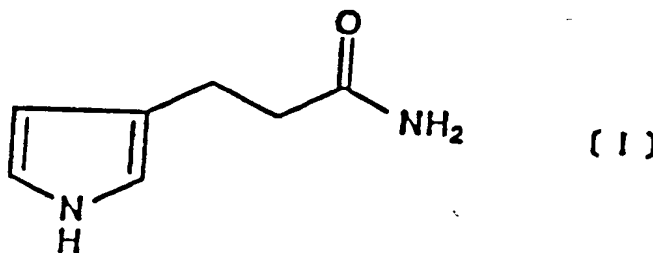
Calpain is a calcium-dependent cysteine protease which occurs in various mammalian tissues. The enzyme is presumed to play a central role in physiological or pathological events, and to be involved in the activation of kinase series enzymes such as protein kinase C and phosphorylase kinase B by limiting proteolysis and in the decomposition of cytoskeletal protein and of growth-factor- and hormone-receptors. Diseases related to calpain have been known. It is strongly suggested to be involved in turnover of muscle in muscular dystrophy, for example, disappearing of Z line in skeletal muscle. (Experimental Medicine, 5 : 942 (1987)).

Furthermore, in a cell infected with human T-cell leukaemia virus, extremely increased activities of calpain and interleukin 2 receptor are observed. This may be caused by irregular reaction of cells to growth factor due to alteration of receptor activity by an action of calpain on cytoskeletal protein. (Biochemistry, 57 : 1202 (1985)). Calpain is also thought to have a relation to myocardial infarction (Experimental Medicine, 5 : 937 (1987)), demyelination (Modern Medicine, 43 : 813 (1988)) and inflammation (ibid., 43 : 776 (1988)).

In a physiological condition, calpastatin, which is a specific inhibitory protein on calpain, is known and is expected to be applied as an effective therapeutic for various excessive calpain-related syndromes. Calpastatin is, however, a high molecular weight-protein and hence it is difficult to apply as a medicine. A low molecular weight-calpain inhibitor is therefore required.

We have found that a microorganism belonging to the genus Streptomyces produces a calpain inhibitor in its cultured broth.

The present invention provides a compound (cystamidin A) of the formula:



or a pharmaceutically or veterinarily acceptable salt thereof.

The present invention also provides a process for the production of the compound of formula (I), or a pharmaceutically or veterinarily acceptable salt thereof, which comprises culturing a microorganism belonging to the genus Streptomyces which is capable of producing the compound of formula (I) in a nutrient medium, isolating the compound of formula (I) from the cultured mass and, if desired, salifying the compound of formula (I) thus obtained.

The compound of formula (I) may be in free form or in the form of a pharmaceutically or veterinarily acceptable salt thereof. Examples of salts are inorganic acid salts such as the hydrochloride and sulfate, and organic acid salts such as the citrate, tartrate or succinate.

The attached drawings show the following:

Fig 1: UV spectrum of cystamidin A;

Fig 2: IR spectrum of cystamidin A;

Fig 3: <sup>1</sup>H-NMR spectrum of cystamidin A; and

Fig 4: <sup>13</sup>C-NMR spectrum of cystamidin A.

Cystamidin A of the present invention can be produced by culturing a cystamidin A producing microorganism belonging to the genus Streptomyces in a nutrient medium, accumulating the cystamidin A in a medium and isolating calpain inhibitor cystamidin A therefrom. The cystamidin A producing microorganism can be a cystamidin A producing microorganism Streptomyces and is not limited a strain illustrated in this specification. Preferable example of cystamidin A producing microorganism is Streptomyces sp. KP-1241 isolated from a soil sample collected in China by the present inventors. Taxonomical properties of the strain KP-1241 is illustrated hereinbelow.

#### 1. Morphological properties:

The vegetative mycelia grow abundantly on both synthetic and complex agar media, and do not show fragmentation into coccoid forms or bacillary elements. The aerial mycelia grow abundantly on inorganic salts-starch agar and glycerol-asparagine agar to show grayish color. The mature sporophores form spiral spore

chains and have more than 20 spores per chain. The spores are oval in shape,  $1.0\ \mu\text{m} \times 0.7\ \mu\text{m}$  in size and have a spiny surface. Whirls, sclerotic granules, sporangia and flagellated spores are not observed.

## 5 2. Cultural characteristics:

The cultural characteristics are shown in Table 1. To investigate the cultural characteristics and physiological properties of the strain, the International Streptomyces Project (ISP) media recommended by Shirling and Gottlieb [Int. J. Syst. Bacteriol., 16 : 313-340. (1966) ] was used. Color Harmony Manual, 4th Ed., 1958 (Container Corporation of America, Chicago) was used for color names and hue numbers. Cultures were observed after incubation at 27 °C for two weeks, if not specified.

15 Table 1

### Cultural characteristics of strain KP-1241

20 Medium	Cultural characteristics
25 Sucrose-nitrate agar	G: Poor, colorless R: Silver gray (3fe) AM: Poor, silver gray (3fe) SP: None
35 Glucose-asparagine agar (ISP)	G: Good, light ivory (2ca) R: Light wheat (2ea) AM: Abundant, silver gray (3fe) SP: None
45 Glycerol-asparagine agar (ISP)	G: Good, colorless R: Bamboo (2gc) AM: Abundant, silver gray (3fe) SP: None

Continued Table 1

<p>5</p> <p>10</p> <p>Inorganic salts-starch agar (ISP)</p>	<p>G: Good, light ivory (2ca)</p> <p>R: Covert tan (2ge)</p> <p>AM: Abundant, dark covert gray (2ih)</p> <p>SP: None</p>
<p>15</p> <p>20</p> <p>Tyrosine agar (ISP)</p>	<p>G: Moderate, light ivory (2ca)</p> <p>R: Pale pink (3ca)</p> <p>AM: Poor, white (a)</p> <p>SP: Poor</p>
<p>25</p> <p>30</p> <p>Oatmeal agar (ISP)</p>	<p>G: Moderate, colorless</p> <p>R: Silver gray (3fe)</p> <p>AM: Moderate, beige gray (3ih)</p> <p>SP: None</p>
<p>35</p> <p>40</p> <p>Yeast extract-malt extract agar (ISP)</p>	<p>G: Good, colorless</p> <p>R: Mustard (2le)</p> <p>AM: Abundant, dark covert gray (2ih)</p> <p>SP: None</p>
<p>45</p> <p>50</p> <p>Nutrient agar</p>	<p>G: Good, colorless</p> <p>R: Bisque (3ec)</p> <p>AM: Moderate, white (a)</p> <p>SP: None</p>

Continued Table 1

Peptone-yeast extract iron agar (ISP)	G: Moderate, light ivory (2ca) R: Honey gold (2ic) AM: Moderate, white (a) SP: None
Glucose-nitrate agar	G: Moderate, light mustard tan(2ie) R: Mustard (2le) AM: Poor, white (a) SP: None
Glycerol-calcium malate agar	G: Good, light ivory (2ca) R: Light ivory (2ca) AM: Poor, white (a) SP: None
Glucose-peptone agar	G: Moderate, bamboo (2gc) R: Bisque (3ec) AM: Moderate, white (a) SP: Amber (3pe)

Abbreviations: G; growth of vegetative mycelium

R; reverse

AM; aerial mycelium

SP; soluble pigment

ISP; International Streptomyces Project

## 3. Physiological properties:

		(+ : Active, - : inactive)
(1)	Melanin formation	
(a)	Tyrosine agar	+
(b)	Peptone-yeast extract iron agar	+
(c)	Glucose-peptone-gelatin medium (21-23°C)	-
(d)	Tryptone-yeast liq.	-
(2)	Tyrosinase reaction	+
(3)	H <sub>2</sub> S production	-
(4)	Nitrate reduction	+
(5)	Liquefaction of gelatin (21-23°C)	+
(6)	Hydrolysis of starch	+
(7)	Coagulation of milk (27°C)	-
(8)	Peptonization of milk (27°C)	+
(9)	Temperature range for growth	12-34°C
	Optimum temperature range for growth	24°C
(10)	Utilization of carbon sources	
	(Pridham and Gottlieb agar medium)	
	Utilized: D-glucose, D-mannitol, D-fructose, i-Inositol	
	Weakly utilized: L-arabinose, D-rhamnose	
	Not utilized: D-xylose, raffinose, melibiose, sucrose	
(11)	Cellulolytic activity	

## 4. Chemical composition:

The DAP (diaminopimelic acid)-isomer in the cell wall of strain KP-1241 is determined to be LL-type.

Taxonomical properties of the strain KP-1241 are illustratively summarized as follows. The DAP-isomer in the cell wall is LL-type. The aerial mycelia form spiral spore chains with smooth surface. In the cultural characteristics, the vegetative mycelia show white or gray on various media. Soluble pigment is slightly produced in tyrosine agar and glucose-peptone agar.

From the taxonomic properties described above, strain KP-1241 is considered to the white or gray series of the genus Streptomyces according to the classification by Pridham and Tresner (Bergey's Manual of Determinative Bacteriology, Vol. 8, pp. 748-829. 1974). The strain is referred to genus Streptomyces and is designated as Streptomyces sp. KP-1241. The strain was deposited in National Institute of Bioscience and Human-Technology, Agency of Industrial Science and Technology, Japan, under the name Streptomyces sp. KP-1241 and the accession No. is FERM BP-4171.

In the present invention, the above strain is merely illustrative, and the above strain, its mutant and the cystamidin A producing microorganism belonging to genus Streptomyces can be naturally used. The calpain inhibitor, cystamidin A of the present invention can be produced by inoculating and culturing aerobically the above strain, for example, in a nutrient medium suitable for cystamidin A production. Nutrient source such as nutrient medium for Streptomyces can be used.

Commercially available nitrogen sources such as peptone, meat extract, corn steep liquor, cotton seed

meal, peanut meal, soybean meal, yeast extract, NZ-amine, casein hydrolysate, sodium nitrate, ammonium nitrate and ammonium sulfate, and carbon sources such as glycerin, starch, glucose, galactose, lactose and mannose can be used. Further, carbon sources such as fatty substance, and inorganic salts such as sodium chloride, phosphate salts, calcium carbonate and magnesium sulfate can also be used.

The other essential trace metallic salt and anti-foam agent such as animal, vegetable or mineral oil can be added to the medium if required. These nutrient sources and other additives useful for production of cystamidin A can be used and that is to say any cultural material for *Streptomyces* can preferably be used in the present invention. Liquid culture such as submerged aeration culture is preferable for mass production of cystamidin A. Temperature for culturing the microorganisms can be adjusted to grow the strain within a range of suitable production of cystamidin A. Culturing condition can be selected and controlled for production of cystamidin A in which depending upon the nature of the above production microorganisms.

Cystamidin A is mainly produced in a cultured broth. A crude substance of cystamidin A can be isolated by extracting the cultured filtrate with water immiscible organic solvent such as butanol, or by eluting the absorbed substance in an absorption resin with water-containing organic solvent. In addition to the above extraction procedure, the isolation method applied for low molecular weight substance such as absorption chromatography, gel-filtration chromatography, preparative thin layer chromatography, counter current partition chromatography and high performance liquid chromatography, and preferable combination and repeating operation thereof, can be applied for isolating and obtaining pure cystamidin A.

Physico-chemical properties of cystamidin A are shown in the followings.

(1) Nature : white powder

(2) Molecular weight : 138.0793 (by mass spectrometric analysis)

(3) Molecular formula:  $C_7H_{10}N_2O$

(4) Melting point : 146 - 148 °C

(5) Ultraviolet absorption maximum (in methanol) : (Fig.1)

203 nm ( $\epsilon = 4400$ ), 213 nm (shoulder,  $\epsilon = 4100$ ), 260 - 280 nm (shoulder,  $\epsilon = 300$ )

(6) Infrared absorption maximum (KBr tablet) : (Fig. 2)

3400, 3250, 3150, 1640, 1570, 1400, 1280, 1110, 970, 805, 740  $cm^{-1}$

(7)  $^1H$ -NMR spectrum (in DMSO- $d_6$ ) : (Fig. 3)(ppm)

10.40 br. s (1H), 7.24 br. s (1H), 6.69 br. s (1H), 6.60 dd (1H), 6.50 dd (1H), 5.86 dd (1H), 2.59 t (2H), 2.25 t (2H) ppm

(s; singlet, d; doublet, t; triplet, br; broad)

(8)  $^{13}C$ -NMR spectrum (in DMSO- $d_6$ ) : (Fig. 4)(ppm)

174.3 s, 121.8 s, 117.3 d, 114.6 d, 107.5 d, 37.0 t, 22.8 t

(9) Solubility in solvent : soluble in water, methanol, acetone, chloroform and benzene

(10) Color reaction : positive for ninhydrin, Ehrlich and Rydon-Smith reagents

From the above physico-chemical properties and the spectra data, the structure of cystamidin A was elucidated as shown in the formula [I] hereinbefore.

As illustrated in details on the physico-chemical properties of cystamidin A, the substance has never been known and reported and is a novel compound.

Biological properties of cystamidin A are illustrated as follows.

Inhibitory activities of cystamidin A against calpain using casein as substrate according to the method by Saito et al. [Agr. Biol. Chem., 51 : 361 (1987) ],  $IC_{50}$  (50 % inhibition) is 0.20  $\mu g/ml$ .

Cystamidin A shows no antimicrobial activities at 100  $\mu g/disk$  (paper disk method) against various kinds of bacteria, yeast and fungi.  $IC_{50}$  value against B-16 melanoma cells in vitro were more than 25  $\mu g/ml$ . The  $LD_{50}$  (ip) of cystamidin A in mice are  $>200 mg/kg$ .

#### Effect of the Invention:

As explained hereinaboves, the novel compound cystamidin A is a low molecular weight substance and shows strong inhibitory action on calpain, therefore it can be used not only for reagent but also for pharmaceuticals.

#### Examples:

The embodiments of the present invention will be illustratively explained in the following examples, but are not construed as limiting.



## Example 1

A loopful mycelia of a stock culture of *Streptomyces* sp. KP-1241 FERM BP-4171 was transferred into 14 flasks consisting of a 500-ml Erlenmeyer flask containing 100 ml of a medium consisting of glucose 0.1%, soluble starch (Kanto Chem. Co.) 2.4%, peptone (Kyokuto Pharmaceutical Industrial Co.) 0.3%, meat extract (Kyokuto Pharmaceutical Industrial Co.) 0.3%, yeast extract (Oriental Yeast Co.) 0.5% and  $\text{CaCO}_3$  0.4% (pH 7.0 before sterilization), and then incubated on a rotary shaker at 27 °C for 3 days to give a seed culture. The seed culture (1.4 lit.) was inoculated into a tank fermenter containing 70 liters of a medium consisting of glycerol 2.0%, soybean meal 2.0% and NaCl 0.3% (pH 7.0 before sterilization), and incubated with agitation at 200 rpm and aeration at the rate of 35 lit./min. at 27°C for 4 days.

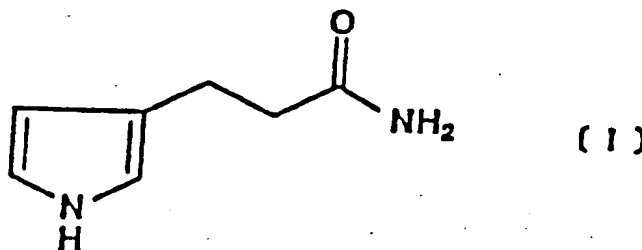
The cultured broth was centrifuged and the supernatant was charged on a column of Diaion HP20 (5 lit., Mitsubishi Chem. Ind. Ltd.) and eluted with 20% methanol. Eluate (25 lit.) containing calpain inhibitory activity was extracted twice with ethyl acetate (18 lit.). The extract was concentrated in vacuo to obtain brownish oil I (55g). The oil was dissolved in water (1.5 lit.), put on a reversed phase silica gel column (1000 ml. YMC \* GEL ODS-AQ 120-S50. YMC Co.) packed with 50% methanol, and developed with 20% methanol.

Active fractions were collected and concentrated in vacuo to obtain brownish oil II (7.2 g). The oil was dissolved in a small volume of methanol, and the solution was mixed with silica gel powder (50 g, Merck Art. 7734), then methanol was removed from the mixture under reduced pressure. The dried silica gel was put on a column of silica gel (210 g, Merck Art 7734) and eluted with a mixture of chloroform and methanol (30 : 1). Active fraction was collected and evaporated under reduced pressure to give brownish oil III (439 mg).

The oil was dissolved in a small volume of methanol and applied on HPLC (Capcell Pak C18, SG120,  $\phi$  20 x 250 mm, Shiseido Co.) separately with 12 times, with 5% acetonitrile at a flow rate of 8 ml/min. by checking and detecting with absorption at 205 nm. Peak showing calpain inhibitory activity, eluted at 17 min. was collected. The extract was concentrated under reduced pressure to obtain white powder of cystamidin A (370 mg).

## Claims

1. A compound of formula:



or a pharmaceutically or veterinarily acceptable salt thereof.

2. A process for the production of a compound of formula (I), or a pharmaceutically or veterinarily acceptable salt thereof, according to claim 1 which comprises culturing a microorganism belonging to the genus *Streptomyces* which is capable of producing the compound of formula (I), in a nutrient medium, isolating the compound of formula (I) from the cultured mass and, if desired, salifying the compound of formula (I) thus obtained.
3. A process according to claim 2 wherein the microorganism is *Streptomyces* sp. KP-1241 (FERM BP-4171) or a mutant thereof which is capable of producing a compound of formula (I) as depicted in claim 1.
4. A compound of formula (I), or a pharmaceutically or veterinarily acceptable salt thereof, as depicted in claim 1 for use in a method of treatment of the human or animal body by therapy.
5. A compound of formula (I), or a pharmaceutically or veterinarily acceptable salt thereof, as depicted in claim 1 for use as a calpain inhibitor in a method of treatment of the human or animal body by therapy.
6. Use of a compound of formula (I), or a pharmaceutically or veterinarily acceptable salt thereof, as depicted

in claim 1 in the manufacture of a medicament for use as a calpain inhibitor.

7. A pharmaceutical or veterinary composition which comprises a compound of formula (I), or a pharmaceutically or veterinarily acceptable salt thereof, as depicted in claim 1 and a pharmaceutically or veterinarily acceptable diluent or excipient.
8. Streptomyces sp. KP-1241 (FERM BP-4171) or a mutant thereof which is capable of producing a compound of formula (I) as depicted in claim 1.
9. A substantially pure culture of Streptomyces sp. KP-1241 (FERM BP-4171), or a mutant thereof which is capable of producing a compound of formula (I) as depicted in claim 1, comprising a source of assimilable carbon, a source of assimilable nitrogen and inorganic salts.
10. A compound which has the following physico-chemical properties:
  - (1) Nature : white powder;
  - (2) Molecular weight : 138.0793 (by mass spectrometric analysis);
  - (3) Molecular formula:  $C_7H_{10}N_2O$ ;
  - (4) Melting point: 146 - 148°C;
  - (5) Ultraviolet absorption maxima (in methanol):  
203 nm ( $\epsilon=4400$ ), 213 nm (shoulder,  $\epsilon=4100$ ), 260 - 280 nm (shoulder,  $\epsilon=300$ );
  - (6) Infrared absorption maxima (KBr tablet):  
3400, 3250, 3150, 1640, 1570, 1400, 1280, 1110, 970, 805, 740  $cm^{-1}$ ;
  - (7) Solubility in solvents: soluble in water, methanol, acetone, chloroform and benzene; and
  - (8) Colour reaction: positive for ninhydrin, Ehrlich and Rydon-Smith reagents;
 or a pharmaceutically or veterinarily acceptable salt thereof.
11. A process for the production of a compound according to claim 10, or a pharmaceutically or veterinarily acceptable salt thereof, which comprises culturing a microorganism belonging to the genus Streptomyces which is capable of producing said compound in a nutrient medium, isolating said compound from the cultured mass, and, if desired, salifying the compound thus obtained.

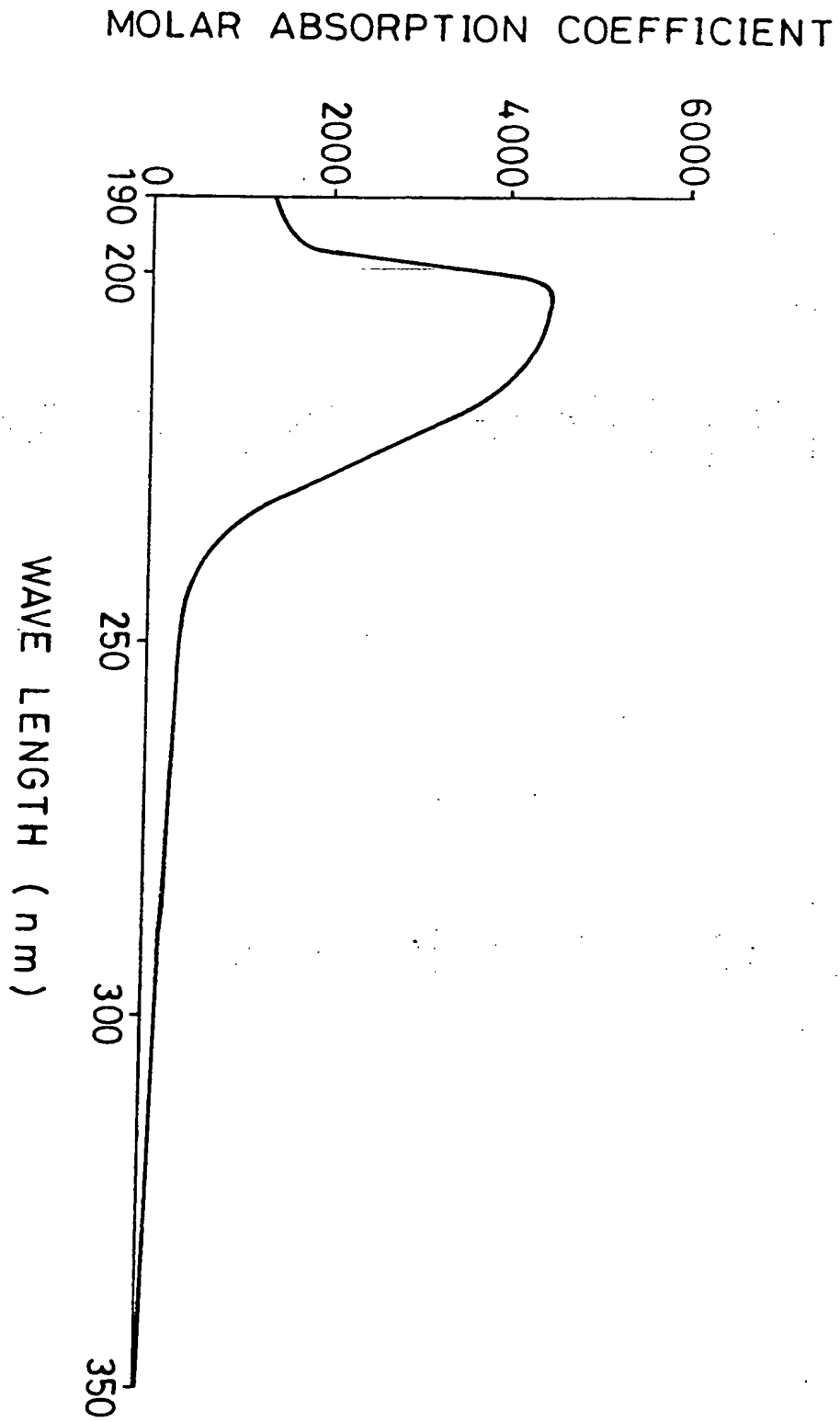


FIG. 1

FIG. 2

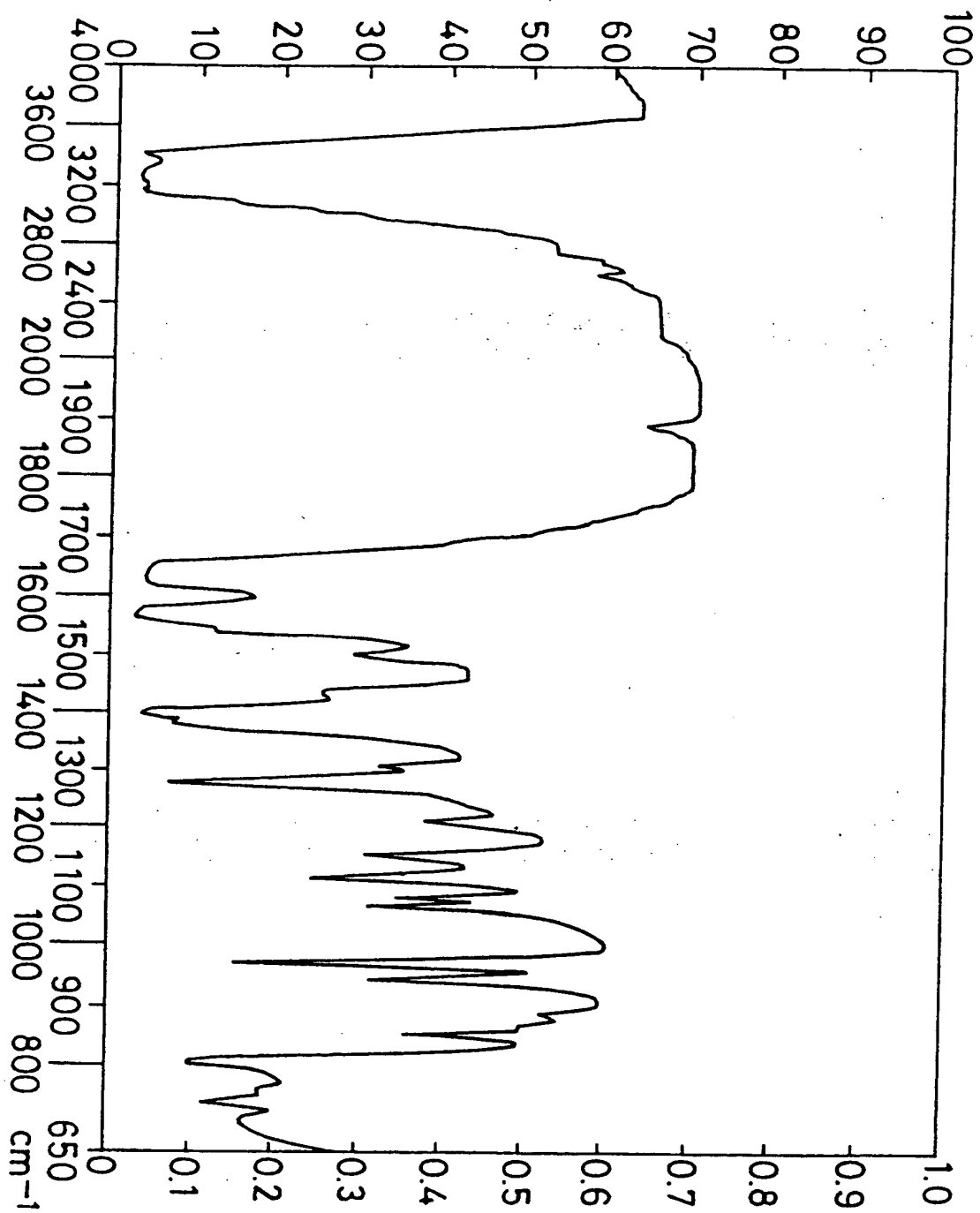
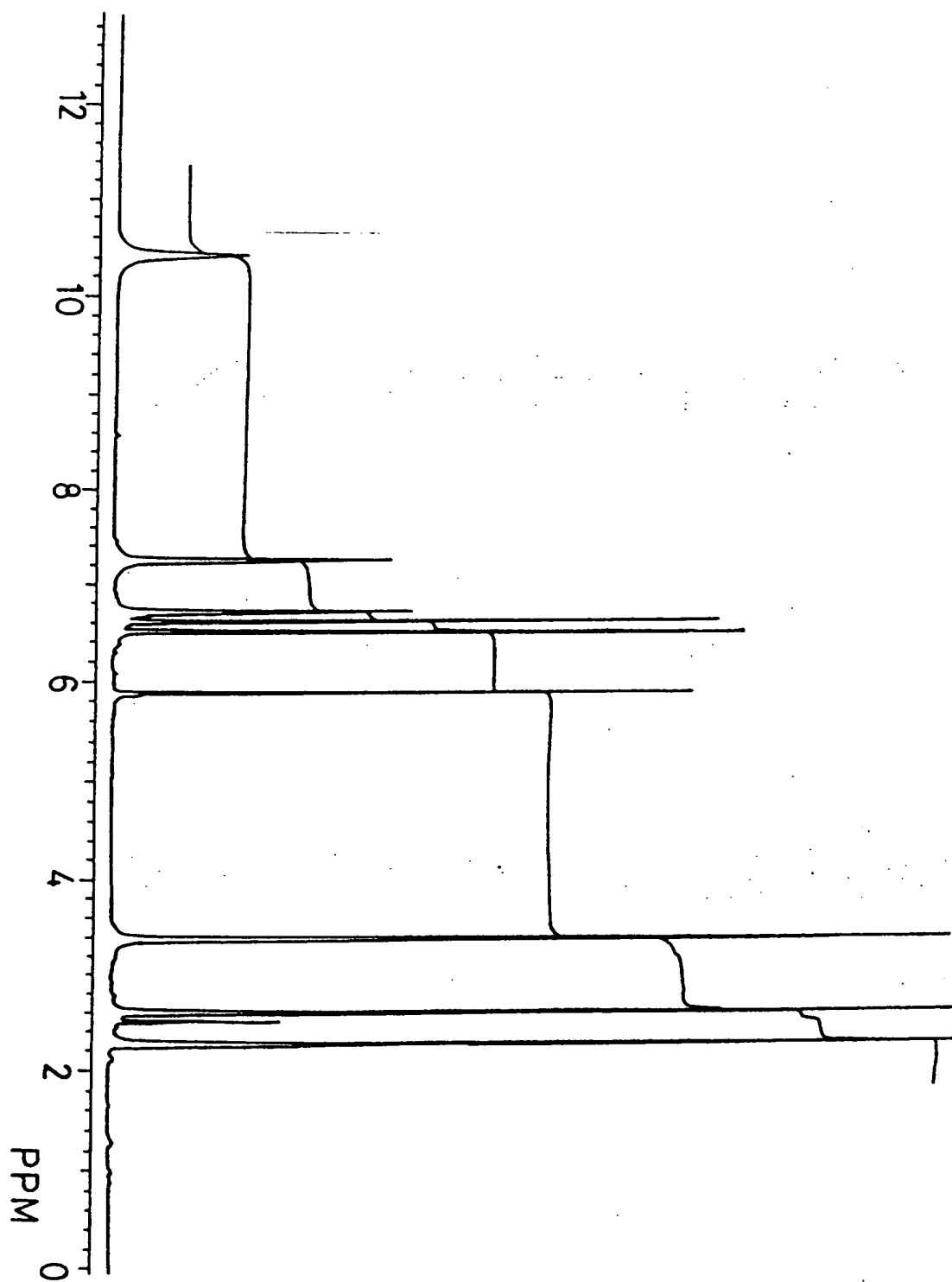


FIG. 3



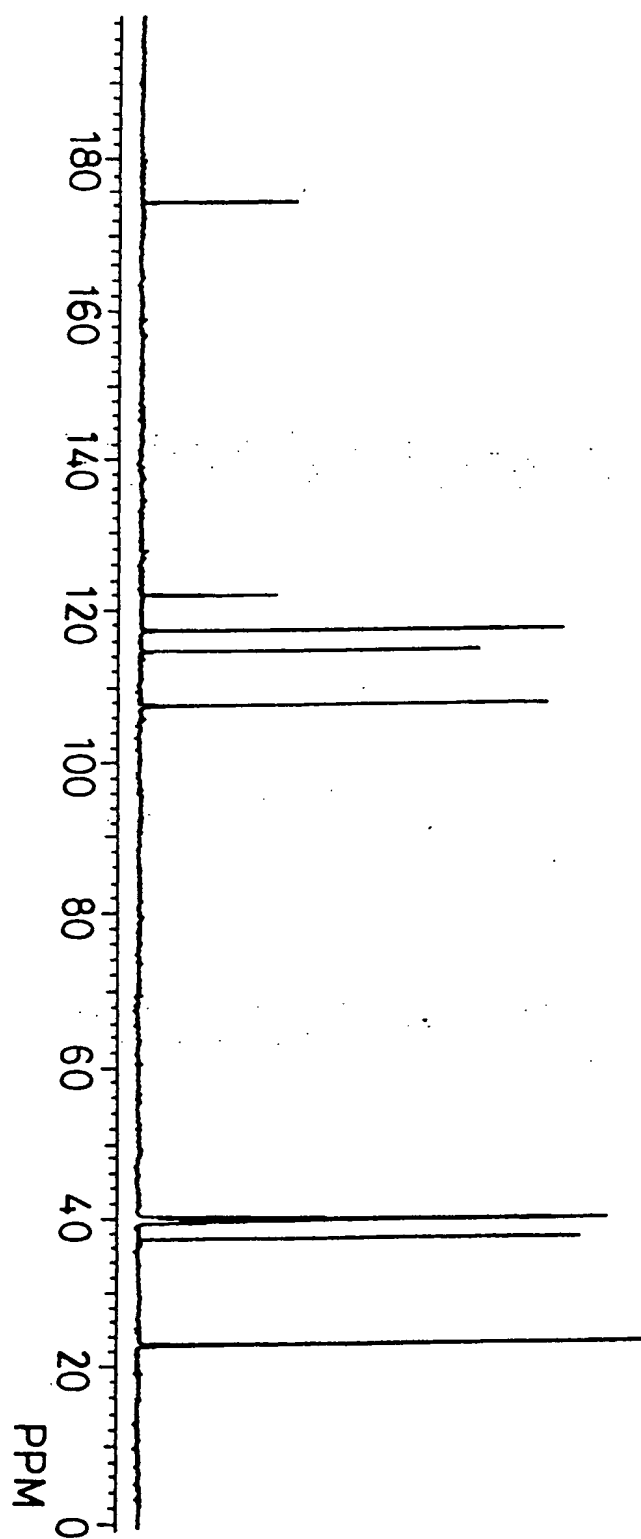


FIG. 4



European Patent  
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# EUROPEAN SEARCH REPORT

Application Number

EP 93 30 2111.5

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
A	US-A-4 376 778 (MEIJI SEIKA KAISHA LTD.) 15 March 1983 * the whole document *	1-5,7-11	C07D207/337 A61K31/40 C12P17/10
A	PATENT ABSTRACTS OF JAPAN vol. 016, no. 329 (C-0963)17 July 1992 & JP-A-04 095 069 ( KIRIN BREWERY CO LTD ) 27 March 1992 * abstract *	1-5,7-11	
A	PATENT ABSTRACTS OF JAPAN vol. 015, no. 388 (C-0872)2 October 1991 & JP-A-03 157 366 ( TAKEDA CHEM IND LTD ) 5 July 1991 * abstract *	1-5,7-11	
A	HOPPE-SEYLER'S ZEITSCHRIFT FÜR PHYSIOLOGISCHE CHEMIE vol. 289, no. 1, 1952, pages 229 - 233 KUTSCHER W & KLAMERTH O 'Darstellung von histaminähnlichen Substanzen aus der Pyrrolreihe.. II. Mitteilung: Die Synthese von 2-[beta-Amino-äthyl]-pyrrol' * the whole document *	1,10	
P,A	JOURNAL OF MEDICINAL CHEMISTRY vol. 35, no. 11, 29 May 1992, pages 2048 - 2054 HUANG Z ET AL. 'Ester and amide derivatives of E64c as inhibitors of platelet calpains' * the whole document *	1,4-7,10	
The present search report has been drawn up for all claims			TECHNICAL FIELDS SEARCHED (Int. Cl.5)  C07D A61K C12P
Place of search MUNICH		Date of completion of the search 06 AUGUST 1993	Examiner HARTRAMPF G.W.
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons A : member of the same patent family, corresponding document	

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